COMPONENTS:

- 1. Manganese (II) sulfite; MnSO₃;
 [13568-71-3]
- 2. Water; H₂O; [7732-18-5]

EVALUATOR:

H.D. Lutz, Dept. of Chemistry, University of Siegen, FR Germany.

April 1983.

CRITICAL EVALUATION:

Manganese sulfite crystallizes from aqueous solutions in the form of several hydrates. The formation of the various hydrates depends on temperature, amount of sulfurous acid, and preparation technique.

Two different trihydrates, monoclinic $\alpha-MnSO_3.3H_2O$ (1-3) [60365-38-0] and orthorhombic $\beta-MnSO_3.3H_2O$ (4) [60365-38-0], and $MnSO_3.5/2H_2O$ (2,5-6) [75042-11-4] are reported to exist at room temperature. At higher temperatures $MnSO_3.2H_2O$ (2) [75042-12-5] and $MnSO_3.H_2O$ (1,7-9) [65410-83-5] are formed. The existence of a $MnSO_3.\frac{1}{2}H_2O$ [60365-38-0] (10) has not been fully verified.

The solubility of these hydrates has not yet been throroughly investigated. Several authors report that manganese sulfite is insoluble in water (7,11-12), readily soluble in sulfurous acid (7) or in other acids, undergoing decomposition (11,12). Only Gorgeu (7) and Rodin et al. (13) give some numerical data, namely for MnSO₃.3H₂O, MnSO₃.2.5H₂O, and MnSO₃.1H₂O. It is not known which of the two different trihydrates Gorgeu investigated, but it seems to be the monoclinic α -MnSO₃.3H₂O (2). The data given by Gorgeu (7) are 0.1 g MnSO₃/kg H₂O (m(MnSO₃) = 7.4 x 10⁻⁴ mol kg⁻¹) at room temperature (cold water) and 0.2 g MnSO₃/kg H₂O (m(MnSO₃) = 1.5 x 10⁻³ mol kg⁻¹) in hot water. The solubility given by Gorgeu seems to be rather too high, because the samples used were probably not fully free of sulfate. Rodin et al. (13) report that the solubility of MnSO₃.2.5H₂O in water increases from 6.93 x 10⁻³ mass % (m(MnSO₃) = 5.13 x 10⁻⁴ mol kg⁻¹) at 293 K to 1.23 x 10⁻² mass % (m(MnSO₃) = 9.13 x 10⁻⁴ mol kg⁻¹) at 343 K. The solubility of MnSO₃.1H₂O is said to be 1.48 x 10⁻² mass % (m(MnSO₃) = 1.10 x 10⁻³ mol kg⁻¹) (13) at 363 K.

TENTATIVE VALUE

The solubility of $MnSO_3$ (i.e. of the hydrates present) in water at room temperature is approximately 5 x 10^{-4} mol kg⁻¹ (molality).

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COMPONENTS:

- 1. Manganese (II) sulfite; $MnSO_3$; [13568-71-3]
- 2. Water; H₂O; [7732-18-5]

EVALUATOR:

H.D. Lutz, Dept. of Chemistry, University of Siegen, FR Germany.

April 1983.

CRITICAL EVALUATION: (continued)

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COMPONENTS:	ORIGINAL MEASUREMENTS:
1. Manganese sulfite; MnSO ₃ ; [13568-71-3]	Rodin, I.V.; Margulis, E.V.
2. Water; H ₂ O; [7732-18-5]	Zh. Neorg. Khim. <u>1983</u> , 28, 258–9; Russ. J. Inorg. Chem. (Eng. Trans1.) <u>1983</u> , 28, 144.
VARIABLES:	PREPARED BY:
Four temperatures: 293 - 363 K	B. Engelen
EXPERIMENTAL VALUES:	
t/°C	MnSO ₃
10 ⁴ mass	_
20 ^b 69.3	5.134
50 ^b 95.4 70 ^b 123.2	7.067
	9.127
90 ^c 148.4	10.994
The solid phase is claimed to be MnSO3.1H2	
	INFORMATION
METHOD APPARATUS/PROCEDURE: Saturation method. Equilibrium was established by stirring the saturated solutions in thermostatically controlled glass tubes. Equilibrium was tested for analytically - 3 hr was reported to be sufficient. Manganese was determined titrimetrically using peroxodisulfate/Ag+.	Not well defined. Not well defined.
	Not given.
	REFERENCES:

COMPONENTS: 1. Manganese sulfite; MnSO ₃ ; [13568-71-3] 2. Carbon dioxide; CO ₂ ; [124-38-9] 3. Sulfur dioxide; SO ₂ ; [7746-09-5] 4. Water; H ₂ O; [7732-18-5]	ORIGINAL MEASUREMENTS: Gorgeu, A. C. R. Hebd. Seances Acad. Sci. 1883, 96, 341-3.
VARIABLES: Two approximate temperatures CO_2 and SO_2 content	PREPARED BY: B. Engelen

EXPERIMENTAL VALUES:

Solubility of manganese sulfite in cold water and hot water, and in cold water saturated with CO_2 or SO_2 is reported.

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Temperature	Solvent	g MnSO ₃ /	$m(MnSO_3)/mo1 kg^{-1a}$
		kg solvent	
cold	pure water	0.1	7.4×10^{-4}
hot	pure water	0.2	1.5×10^{-3}
cold	water saturated	1 b	7.4×10^{-3c}
	with CO ₂		
cold	water saturated	150-170	1.11 - 1.26
	with SO ₂		

- a Calculated by the compiler.
- b Units are in g/dm³ solvent.
- c Units are in mol/dm³ solvent.

AUXILIARY INFORMATION

METHOD APPARATUS/PROCEDURE:	SOURCE AND PURITY OF MATERIALS:
Not given.	MnSO3.3H2O was prepared by precipitation
o .	from an aqueous solution of MnCl ₂ with
	alkaline sulfite, added in small portions.
	The sulfite contained 2-3% of sulfate
	(author).
	ESTIMATED ERROR:
	REFERENCES:
	1